2. (amended) A pharmaceutical preparation according to claim 1, [characterised in that] wherein the solvent comprises the following components: (a) [of] from about 10% to 20% by weight of ethanol; (b) [of] from about 10% to 60% by weight of glycol; and (c) water[, optionally containing additives,] in a volume percentage adding up to 100% by volume.

Claim 3, line 1, delete "characterised in that" and replace with -- wherein --.

Claim 4, line 1, delete "characterised in that" and replace with -- wherein --.

Claim 5, line 1, delete "characterised in that" and replace with -- wherein --; and line 2, delete "of".

Claim 6, line 1, delete "characterised in that" and replace with -- wherein --; and line 4, after the second occurrence of "water", delete "," and after the first occurrence of "water", insert -- , --.

Claim 7, line 2, delete "the mixed" and replace with -- a mixture of the --.

9. (amended) A process [according to claim 7, characterised in that] for the preparation of a solution in accordance with claim 3 [is made by first], comprising dispersing Tamoxifen Citrate in the propylene glycol to form a dispersion, adding the dispersion to the glycerol, [and,] then adding the ethanol component thereto to form a solution, and then adding the water component.

Please add the following new claims:

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